



## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-392]

#### **Bulk Manufacturer of Controlled Substances Application: AMRI Rensselaer, Inc.**

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before [INSERT 60 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER].

**ADDRESS:** Written comments should be sent to: Drug Enforcement Administration,  
Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive,  
Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.33(a), this is notice that on March 15, 2019, AMRI Rensselaer, Inc., 33 Riverside Avenue, Rensselaer, New York 12144-2951 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled Substance	Drug Code	Schedule
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
Amphetamine	1100	II
Lisdexamfetamine	1205	II
Pentobarbital	2270	II
4-Anilino-N-phenethyl-4-piperidine (ANPP)	8333	II
Codeine	9050	II
Oxycodone	9143	II
Hydromorphone	9150	II
Hydrocodone	9193	II
Meperidine	9230	II

Morphine	9300	II
Fentanyl	9801	II

The company plans to manufacture bulk controlled substances for use in product development and for distribution to its customers. In reference to drug codes 7360 (marihuana) and 7370 (tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetics. No other activities for these drug codes are authorized for this registration.

Dated: August 9, 2019.

Neil D. Doherty,

*Acting Assistant Administrator.*

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